

005211 070522Z60

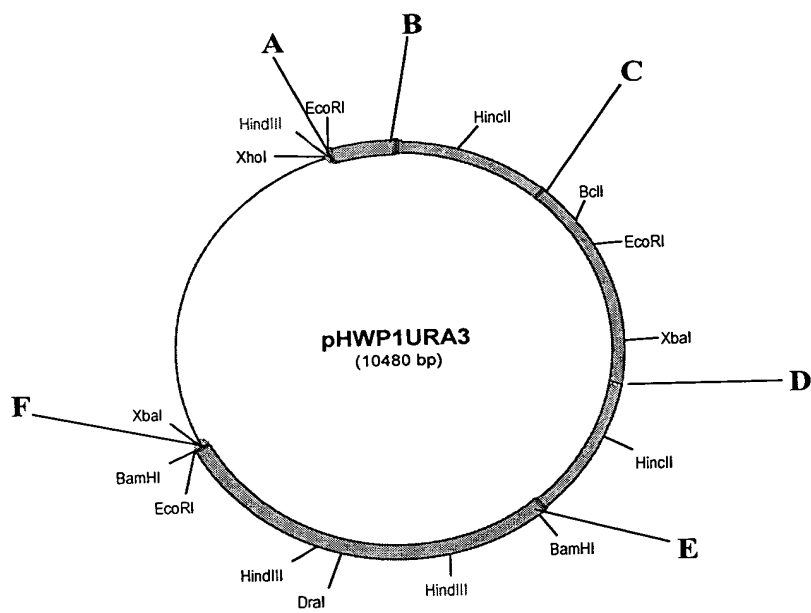


Figure 1

Role of HWP1 in health of mice orally colonized with *C. albicans*.

Mouse type	Health	Number of mice given <i>C. albicans</i> strains of HWP1 type			
		I. <u>HWP1 HETEROZY GOTE</u>	hwp1/hwp1* homozygote	II. <u>HWP1 REVERTANT</u>	Wild type
Beige nude	ill	5	2	3	3
	not ill	2	9	1	1
	total	7	11	4	4
	*P < 0.05 compared to the heterozygote, P = .058 compared to the revertant. P < .05 compared to combined heterozygote and revertant groups. Survival differences between other groups were not significant.				
Epsilon 26	ill	5	0	4	5
	Not ill	0	5	1	0
	total	5	5	5	5
	*P < .01 compared to individually to the heterozygote and revertant. groups. Survival differences between other groups were not significant.				

Figure 2

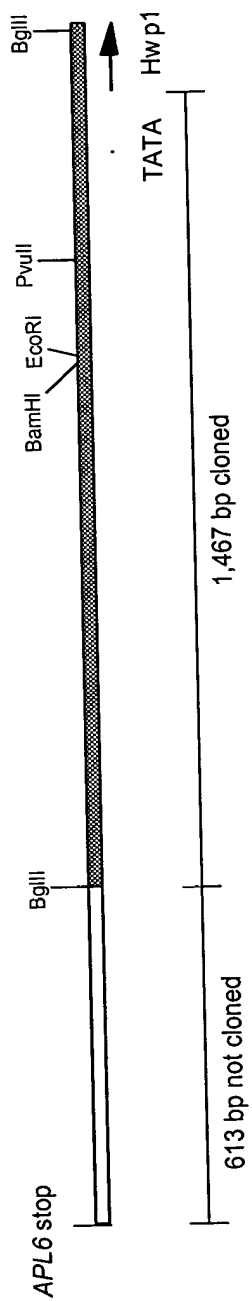


Figure 3

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1  GGATCTTTCTTTTCCCTTAAACCGGATCAAGAAAGAAAGTGGAATAAAGCTATGATAAATGTTGATTTTGTGTAATTC
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171 AGAAATACAGGAAACCCCTCCAAAAAATTTTGGACCTTACACGCACATAAATTTGCGGATAAACTTGCCATAATAAAAACTCT
256 TTGAAACATACGATATGTTATTCTTTTCATAAAGTGAATATTTTGGCTTTTAAACATTATGAACAATTTGAAAAAAGGAA
341 ATGAAAAAGGTAAAGATTGCCTAACCAATTGAAAAATAAAGGCTAAGGTTTCTGATGCGTTTAACTAAAAAGGAAATACAAAA
426 GTTATTAGCGATAACCTGCGTAAGGTGCAACAAAATATATTTGCAGCTTAGCTCTATAGAAAAATATACAACTAAATCCTTAA
511 GGAATTTCCCTCTATATATAATAGGAAATCCCTCTCACAGTGAAGTGAATATCCATCTGAATTTATCAGTCCACTAATTTCCATCAA
596 TAAAAATAGATTAGTGTATTGTTCTCTTCAGTACAATTAATACCATTAATGCAATGCTAGCTTATTGTTTCATAATTAGCCATGTTGC
681 ACACCTTAATTCGAACATTAACTGTATCCATATTTTCTTGCTCTCTTTGTTTCTAACAAAAATGTTCCAGAAATTTTTTA
766 AAAAAATATTTGAAAAAACACATAACACTTTGAGTATGATAATATCAACTATTGACTTGTGTTGAAAGTAAAGAAATCAAAATTTTTT
851 TCTAACTCGACTAATGCACCTTACATCAACTGGATGTTATTGTCATCTACTACTATAAGCTCAAAACAAATTTATCTTTCAAAAAATG
936 TTATAATTAAACAAGTCATCTATAATTCTTTGGATCCAAAAAACAAAGGAATTCGGAAATTTCTGACGATAAATGTGCGACTCACAATTC
1021 ATTGTAAAAAGGGAGAGTTTGGTAGGCTCATAAATCGCTTATAATGTACCTCTAAAGTAATCTAAAAACAAACACAACTTTCTAA
1106 AACCTATAATAATAACCCCTAATGGCTCACAAACCGGGATAAGTTAGTTAGCCCCAGCTGTTTTTTTTTGGCCTTATTTTATGACTA
1191 CATTTGTTTCACCTTTTGTGCGACTTTAATACCGTTTTTGGCACTTCTCTTTGTATCACCTGTATCCGCCCTTTTTTAACATAG
1276 CAACTCTGTAAAGTCCCTTTCTTTTCCCACTATTTTATCATTTCTTGAAATATGTAATCAGAATAGTTTTTCAAAAAACTATAAAT
1361 AACGGTCAAAATAACCGGCTATTTTCAATTTCCATTCAACTTGTTTTCTCAACAATATCAAAACAAACAGGAATCTCCTATAGTC
1446 ACTCGCTTTTAGTTTCGTCAATATG

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BamHI EcoRI

PvuII

TATA box

Figure 4

* * ♦ * *

GAT99 CTNCGTKTTPLWRRNPQGQPLCNACGLFLKLHGVVRPLSLKTDVIKKRQR [SEQ ID. NO: 2]
 :::: ::

NIT2 CTNCFQTTPLWRRNPDGQPLCNACGLFLKLHGVVRPLSLKTDVIKKRNR [SEQ ID. NO: 3]

Figure 5

TAATTCTAATAACTGATACTAAGTTTTGTTCCCTTTTTTTGGGATTTCTTTTTTTCTAATTTT
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 GTTTATCTTTTTCTTCTTCTTCTACTTGCTTTGTTTTCTGTTTTCGGTTTCATAGTTGATAGCTT
 TTAATAAATACCCCTTTTTTTTTTACAATAGTTAGTTCTAAGCTTATTCAGTGGTTTAATTGG
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Figure 6

005211 010522/60

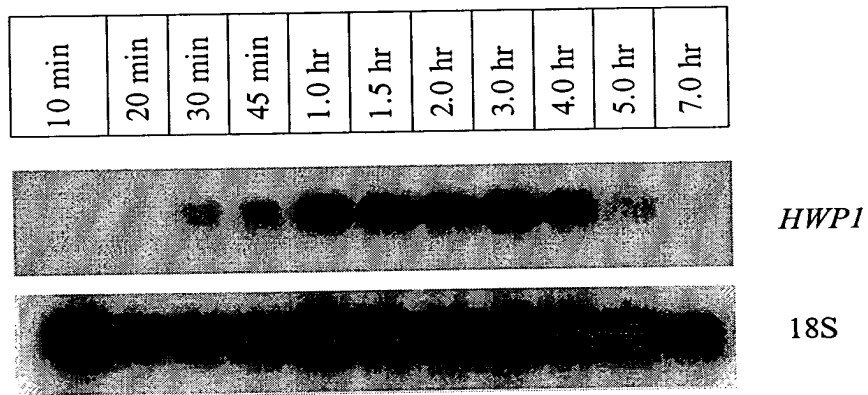


Figure 7

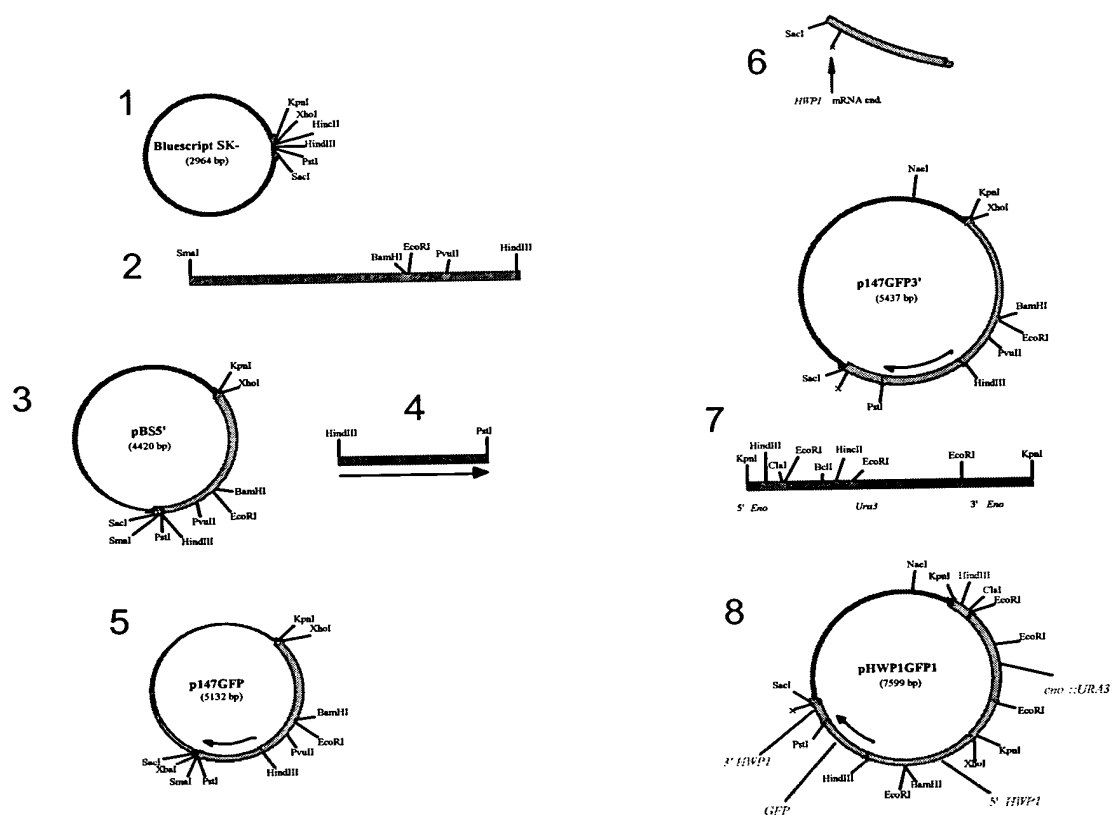


Figure 8

005277 07052260

Integration of pHWP1GFP1 into the chromosome
of *C. albicans* at the enolase locus.

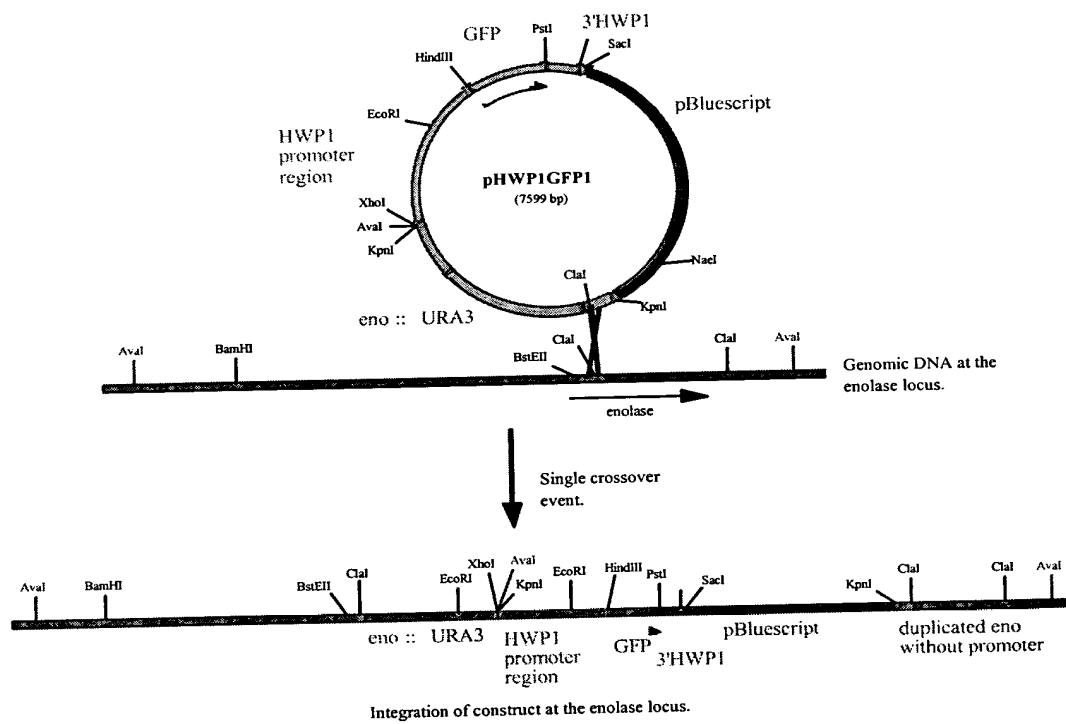


Figure 9

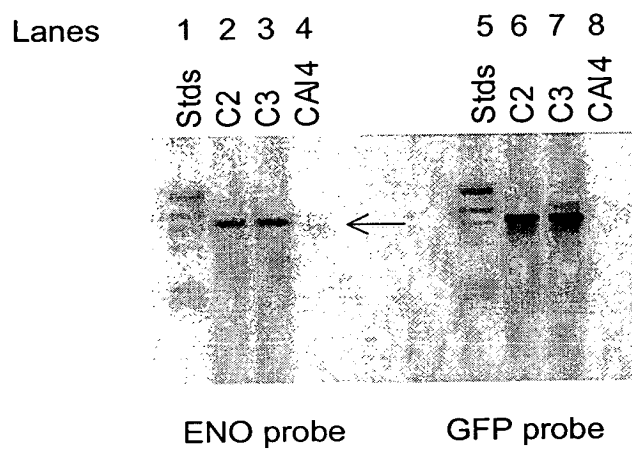


Figure 10

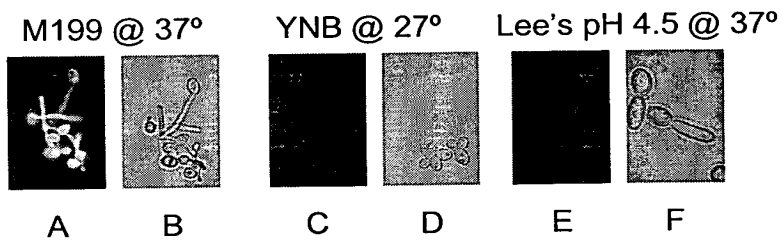
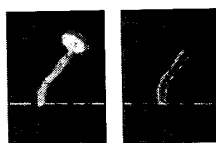


Figure 11

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A B

Figure 12

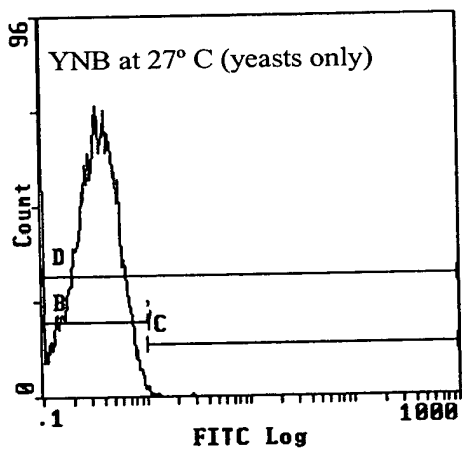


Figure 13A

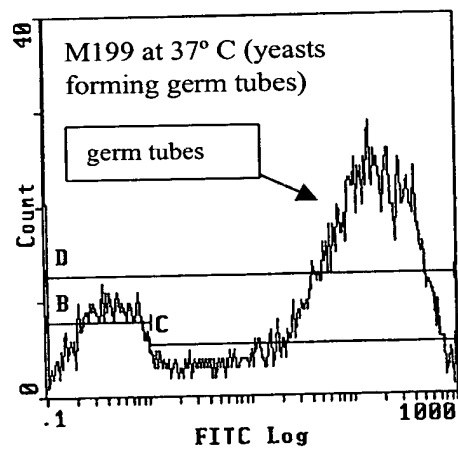
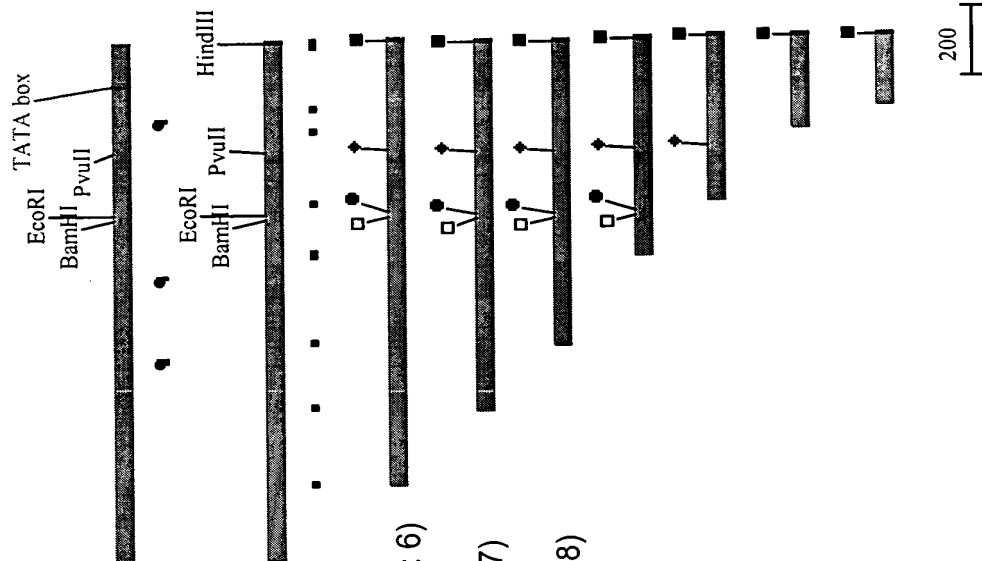


Figure 13B



5' GG CTCGAG GGA CCT TAC ACG CAC ATA AAT TGC Δ205 bp
(SEQ ID. NO: 5)

5' GG CTCGAG CAA AAG TTA TTA GCG ATA ACC TGC Δ421 bp (SEQ ID. NO: 6)

5' GG CTCGAG GTG TAT TGT TCT CTT CAG TAC ATT Δ608 bp (SEQ ID. NO: 7)

5' GG CTCGAG CTC GAC TAA TCG ACT TTA CAT CAA Δ856 bp (SEQ ID. NO: 8)

5' GG CTCGAG ATG TCG ACT CAC AAT TCA TTG Δ1004 bp (SEQ ID. NO: 9)

5' GG CTCGAG GTT GCG ACT TTA ATA CCG TT Δ1209 bp (SEQ ID. NO: 10)

5' GG CTCGAG CAT AGC AAC TCT TGT AAA GTC Δ1271 bp (SEQ ID. NO: 11)

Figure 14A

Figure 14B

Identification of virulence and morphogenesis factors in *C. albicans*

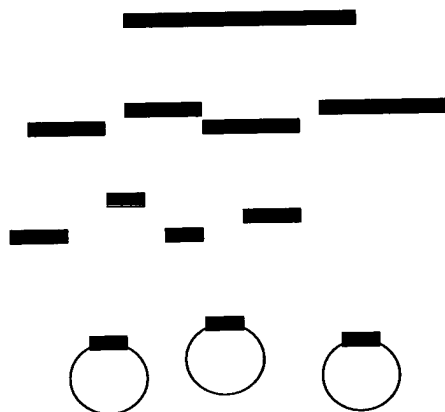
I. STEP 1: CREATE A GENOMIC LIBRARY FOR MICROARRAY CONSTRUCTION

Prepare *C. albicans* genomic DNA.

↓
Sau 3A partial digest.

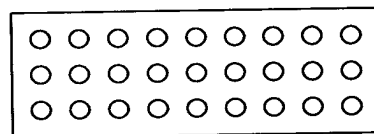
↓
Size selection of 0.5 to 2.0 kb fragments.

↓
Clone genomic fragments into plasmid vector (pBluescript).



A. STEP 2: Create Microarray

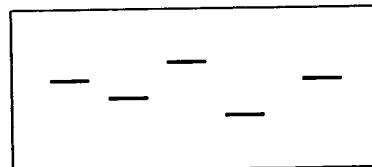
Transfer transformants to 96-well plates.



↓
Perform colony PCR using universal primers.



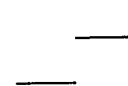
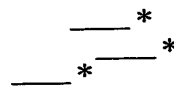
↓
Check PCR rxns on gels and rearray positives on 96-well



↓
Spot productive rxns on membranes.

↓
Prepare and label cDNA from mRNA of strains with and without DNABPG

Labeled cDNA from strain; with DNABPG dnabpg null mutant

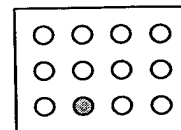
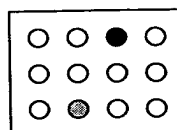


Labeled cDNA.

↓
Hybridize labeled cDNA to duplicate membranes.

wild type

mutant



↓
Go back to 96-well plates and sequence the clones of interest.

→ In vivo analysis of genes.

Figure 15